International application No.

PCT/JP2007/061850

A. CLASSIFICATION OF SUBJECT

A61K45/00(2006.01)i, A61K39/395(2006.01)i, A61P7/06(2006.01)i, A61P35/02 (2006.01)i, A61P43/00(2006.01)i

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols) A61K45/00, A61K39/395, A61P7/06, A61P35/02, A61P43/00

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
Jitsuyo Shinan Koho 1922-1996 Jitsuyo Shinan Toroku Koho 1996-2007
Kokai Jitsuyo Shinan Koho 1971-2007 Toroku Jitsuyo Shinan Koho 1994-2007

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) BIOSIS(STN), CAplus(STN), EMBASE(STN), MEDLINE(STN), WPI(DIALOG), JMEDPlus(JDream2), JST7580(JDream2), JSTPlus(JDream2)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Further documents are listed in the continuation of Box C.

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	JP 10-510842 A (ZymoGenetics, Inc.), 20 October, 1998 (20.10.98), Claims; page 8, lines 21 to 24; examples & WO 96/40218 A1 & AU 199662500 A & EP 831888 A1 & US 6013067 A & MX 199709244 A1 & KR 1999022420 A & CN 1190348 A	1-11,18,19, 21-23
х	JP 2005-539082 A (Ortho-Mcneil Pharmaceutical Inc.), 22 December, 2005 (22.12.05), Full text & WO 2004/026332 A1 & AU 2003275077 A1 & EP 1542714 A1 & NO 200501841 A & BR 200314591 A & US 2005/0282277 A1 & CN 1723036 A & ZA 200503036 A & KR 2005093759 A	1-11,18-20, 22,23

* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means document published prior to the international filing date but later than the priority date claimed	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family
Date of the actual completion of the international search 24 July, 2007 (24.07.07)	Date of mailing of the international search report 07 August, 2007 (07.08.07)
Name and mailing address of the ISA/ Japanese Patent Office	Authorized officer
Facsimile No.	Telephone No.

See patent family annex.

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		PC1/JP2(007/061850		
C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT					
Category*	Citation of document, with indication, where appropriate, of the relevant passages		Relevant to claim No.		
X Y	NAKAMURA, T., et al., A Novel Non-Peptidyl C-Mpl Agonist, NIP-004, Stimulates Human Megakaryopoiesis and Thrombopoiesis [onling Blood (ASH Annual Meeting Abstracts), 2005 Abstract 3148. Retrieved from the Internet <url:http: 11="" 3148?maxtoshits="10&hits=10&RESULTFORMAT=&fulltext=NIF" abstract="" abstracts.hematologylibrary.orcontent="" ashmtg;106="" resourcetype="HWCIT" searchid="1&FIRSTINDEX=0&volume=106&issue=1">, full text</url:http:>	ne]. 5, 106: 5: cg/cgi/ show=& 2-004&	1,18 2-11,19-23		
X Y	SUZUKI, K., et al., YM477, a Novel Orally-Thrombopoietin Receptor Agonist [online]. (ASH Annual Meeting Abstracts), 2005, 106: Abstract 2298. Retrieved from the Internet <url:http: 11="" 2298?maxtoshits="10&hits=10&RESULTFORMAT=&fulltext=YM4" abstract="" abstracts.hematologylibrary.orcontent="" ashmtg;106="" resourcetype="HWCIT" searchid="1&FIRSTINDEX=0&volume=106&issue=1">, full text</url:http:>	Blood :: :g/cgi/ :Bhow=& :77&	1,18 2-11,19-23		
Y	WO 2005/107784 Al (Chugai Pharmaceutical Ltd.), 17 November, 2005 (17.11.05), Claims; abstract (Family: none)	Co.,	1-11,18-23		
Y	WO 2005/056604 A1 (Chugai Pharmaceutical Ltd.), 23 June, 2005 (23.06.05), Claims; abstract & EP 1616881 A1 & NO 200603224 A & US 2006/0222643 A1 & AU 2004297111 A1 & MX 2006006289 A1 & BR 200417077 A & KR 2006107572 A		1-11,18-23		
Y	WO 02/33072 A1 (Chugai Pharmaceutical Co. Ltd.), 25 April, 2002 (25.04.02), Claims; abstract & AU 200210917 A & EP 1327680 A1 & KR 2003055274 A & CN 1469925 A & US 2004/0091475 A1 & CN 1721445 A	,	1-11,18-23		
A	JP 2004-222502 A (Asahi Kasei Corp.), 12 August, 2004 (12.08.04), (Family: none)		1-11,18-23		
A	JP 2005-204539 A (Mitsubishi Pharma Corp. 04 August, 2005 (04.08.05), & WO 2005/071064 A1),	1-11,18-23		

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Box No. II	Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)
1. X Cla bec The i an agor for tr in the 2. Cla bec.	onal search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons: tims Nos.: 12-17 ause they relate to subject matter not required to be searched by this Authority, namely: nvention according to claims 12-17 includes the step of administering nist for TPO receptor to a test subject, and corresponds to a method eatment of the human body by therapy because the human is included test subject. (PCT Article 17(2)(a)(i), PCT Rule 39.1(iv)) tims Nos.: ause they relate to parts of the international application that do not comply with the prescribed requirements to such an ent that no meaningful international search can be carried out, specifically:
	ims Nos.: ause they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box No. III	Observations where unity of invention is lacking (Continuation of item 3 of first sheet)
1. \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	all required additional search fees were timely paid by the applicant, this international search report covers all searchable
clai	ms.
	all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of additional fee.
	only some of the required additional search fees were timely paid by the applicant, this international search report covers y those claims for which fees were paid, specifically claims Nos.:
	required additional search fees were timely paid by the applicant. Consequently, this international search report is ricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remark on the	Protest The additional search fees were accompanied by the applicant's protest and, where applicable, payment of a protest fee
	The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
	No protest accompanied the payment of additional search fees.

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Regarding claims 1-11, 18-23

Claims 1-11, 18-23 describe an agonist for TPO receptor as an active ingredient of a drug or an ingredient to be used for manufacturing a drug.

As the agonist for TPO receptor, a lot of compounds other than TPO can be included, however, even a person skilled in the art cannot clearly realize the whole compounds. Further, even if the description of the specification is examined, it is not described to such an extent that the whole compounds are made clear.

Thus, a sufficient support by the specification is not made by the description of the claims. Further, it is not clearly and sufficiently disclosed to such an extent that a person skilled in the art can make implementation. (PCT Articles 5 and 6)

Incidentally, in the creation of this international search report, a prior art search was carried out based on the description of the specification within a reasonable range.